Stem Cell Clinical Trials: Manufacturing and Regulatory Issues | Keith Wells

Developing an idea for a therapy from the lab bench all the way through approval by the Food and Drug Administration is an extremely challenging and complex process. Keith Wells, Senior Consultant for Biologics Consulting Group, spoke to CIRM-funded stem cell scientists to help them anticipate the manufacturing and regulatory issues that can crop up while developing a stem cell based product for patients.

Learn about California's progress toward stem cell therapies